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(54) Enzyme detergent composition

(57) A detergent composition comprising an alkali cellulase is particularly effective for removing solid, inorganic dirt and enhances the deterging effect of phosphorus free or low phosphorus content detergents. The cellulase may be obtained from *Bacillus* or *Aeromonas* species.

GB 2 095 275 A

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The present invention relates to a detergent composition. More particularly, the present invention relates to a detergent composition characterised by containing an alkali cellulase which exhibits high 5 activity under alkaline conditions.

Recently, techniques of cleansing clothes have been remarkably developed. The washing of clothes has been facilitated greatly by the development of detergent materials, water conditioning, improvements in and increases in the spread of washing machines and improvements in fibers. Among them, the improvement of the starting materials for detergents is remarkable. Owing to the 10 improvements in surfactants, builders, dispersants, fluorescent dyes and bleaching agents, the compositions of detergents for clothes have nearly reached the stage of completion. Detergents for clothes are based on the following principles:

(1) A surfactant or builder is adsorbed on the dirt or/and fiber surface to reduce the surface tension between the dirt or/and fiber and water, so that the dirt is physico-chemically separated out of 15 the fiber.

(2) The dirt is dispersed and solubilized by means of a surfactant or inorganic builder.

(3) The dirt is chemically decomposed by an enzyme such as protease.

(4) Coloured stains are bleached with a bleaching agent.

(5) A fluorescent dye is adsorbed on the fiber surface to brighten it.

(6) The precipitation of active detergent ingredients by divalent metal ions is prevented by means 20 20 of a chelating agent.

The fundamental idea of cleansing clothes in the prior art is the incorporation of a component which directly attacks the dirt or a component which enhances the attacking power of said active ingredient in the composition. At present, the effectiveness of the known detergent compositions based 25 on this fundamental idea have almost reached an optimum level and great efforts will be required if further improvement are to be made.

After intensive investigations from viewpoints different from the conventional ideas of the deterging of clothes, the inventors have found that, unexpectedly, quite excellent deterging effects on types of dirt which are generally not concerned with enzymatic activity of an alkali cellulase (one of the 30 cellulase enzymes) are obtained if the alkali cellulase is incorporated in a detergent. The present irvention has been completed on the basis of this finding.

Commercially available or generally well-known cellulases have an optimum pH under acidic or neutral conditions and their use under alkaline conditions has generally been avoided, since their activity has been reduced under such conditions.

The present invention provides a detergent composition characterised by containing an alkali cellulase having an alkaline pH as its optimum pH. According to the present invention, there is provided a detergent composition having a remarkable deterging effect particularly on inorganic types of dirt which are generally not concerned with alkali cellulase activity at all.

As described above, it has been known in the art to use an enzyme as a component of a 40 detergent. As such enzymes, there have been used only those which effectively act on the dirt. More particularly, there have been used only protease for protein stains, amylase for saccharide stains and lipase for oily and fatty stains. Those enzymes attack the stains directly. Though the mechanism of the deterging effects of the alkali cellulase in the present invention have not fully been elucidated, it is known that the effects are not merely based on an expansion of the fibers, unlike phenomena observed 45 when surfactants are used.

The main advantages of the present invention are that the detergent is particularly effective for removing solid inorganic dirt such as fine mud particles which cannot be removed sufficiently with conventional detergents and also for removing other types of dirt such as the stains on the necks and cuffs of clothes and oily stains on clothes and that it is highly useful for enhancing the deterging power of 50 phosphorus-free detergents and detergents having only a low phosphorus content. Phosphate salts have been effective for the removel of fine mud particles that has penetrated into the fibers. However, there is nowadays a desire to reduce the amount of phosphate salts incorporated into detergents because of the problem of eutrophication. The requirement for the use of phosphate-free detergents makes the removal of mud particles extremely difficult. In particular, the removal of mud particles from 55 c tton clothing is quite difficult, as is well known. Further, muddy stains on canvas shoes have proved 55 difficult to remove.

The present invention throws a fresh light on the abov -menti ned problems. According to the present invention, an xcellent deterging power quivalent or superior to that of a weak alkaline powdery detergent containing sufficient phosphate salt can be brained by, for example, (1) applying 60 th present invention to an alkaline detergent containing no phosphates or a small amount of phosphates or (2) applying the present invention to a weakly alkaline, liquid, phosphate-free detergent in the cleaning of muddy stains in cellulose fibers or blended fiber cloths comprising cellulose fibers and other fibers.

Another great advantage of the present invention is that it can be applied to detergents of any

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shape. The alkali cellulase may be incorporated in a composition in the form of any of spray-dried p wder, p wder-blended powder, tablets or liquid to obtain the detergent composition of the present invention.

The alkali cellulases used in the present Invention are those having an optimum pH of 8.0—11.5, 5 preferably 8.1—11.0. Under alkaline conditions, those enzymes exhibit an activity equivalent to that exhibited under acidic or neutral conditions. The alkali cellulases are obtained from cellulase culture liquids of various origins by refining and fractionation according to salting out, precipitation, dialysis or gel permeation method. The alkali cellulases are obtained also by fractionation of crude enzymes or refined enzymes according to electrophoresis or by the heat treatment of the same (for example, at 10 40-90°C for 0.5 min to 3h).

As particularly preferred alkali cellulases used in the present invention, the following enzymes

may be mentioned:

The specific cellulase to be used in the invention is preferably a cellulase produced by a fungus of Bacillus N or a cellulase 212-producing fungus belonging to the genus Aeromonas. The Bacillus N is disclosed in Japanese patent publication No. 28515/75 and available from the Fermentation Research Institute, the Agency of Industrial Science and Technology, located at 1-1-3, Higashi, Tsukuba-Yatabemachi, Ibaraki, Japan. It has been added to its permanent collection of micro-organisms as deposition numbers, FERM Nos. 1138, 1139, 1140 and 1141. The Aeromonas fungus has been added also to the permanent collection of microorganism in the Fermentation Research Institute, the Agency of Industrial Science and Technology, as FERM No. 2306 and is disclosed in Japanese patent publication (unexamined) No. 76287/75, now published after examination as Japanese patent publication

The cellulase to be used in the invention includes a cellulase extracted from the hepatopancreas of a marine mollusc (Dolabella Auricula Solander), which is disclosed in Biochem. J. (1966) 99, 214—

25 221.

Each of celluloses produced by these fungi is a special cellulase which retains a high activity even under alkaline conditions and has an alkali resistance.

The detergent composition of the present invention is characterized in that this special cellulase is contained as one indispensable ingredient. More particularly, the present invention provides a 30 detergent composition having a prominent washing power to inorganic stains irrelevant to the inherent activity of the cellulase, especially collar contaminations consisting of mixtures of inorganic stains and oils secreted on the skin surface, which change with the lapse of time.

When the detergent composition comprising a special cellulase having a high activity under alkaline conditions and also having an alkali resistance, which is produced by a fungus selected from 35 Bacillus N (deposited with FRI deposition numbers of 1138 through 1141) and a cellulase 212producing fungus belonging to the genus Aeromonas, is employed, an excellent washing effect can be obtained over a broad range of the pH value of the washing bath.

This excellent effect overbalances reduction of the washing power due to reduction of the alkali capacity of the builder on reduction of the pH value of the washing bath.

The enzymatic activity of the cellulase that is used in the present invention is determined

according to the following method.

50mg of Avicel (for chromatography) or carboxymethyl cellulose (CMC) is suspended in 4ml of a glycine NaCl-NaOH buffer solution (having a pH value of 8,3), and the suspension is preheated at 37°C for 5 minutes and 1ml of an enzyme liquid is added to the suspension. The suspension is sufficiently mixed and reaction is carried out for 1 hour. After completion of the reaction, the quantity of reduced sugar is determined according to the 3,5-dinitrosalicyclic acid method. More specifically, the liquid reaction mixture is filtered, and 3ml of 3,5-dinitrosalicylic acid is added to 1 ml of the filtrate and the mixture is heated at 100°C for 10 minutes to effect coloration. The mixture is cooled and is then mixed with deionized water so that the total volume is increased to 25ml. The resulting liquid is subjected to 50 colorimetry at a wavelength of $500m\mu$.

When 1 mg of the enzyme as the solid produces reduced sugar in an amount corresponding to 1μ mole of glucose for 1 hour under the above conditions, the enzymatic activity is defined as 1 unit per

mg of the solid.

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The present invention is attained by combining those alkall cellulases with well-known detergent 55 compositions. As for the alkali cellulase content, it is preferred that the composition contains 0.01wt.%, particularly 0.1—10 wt.%, of an alkali cellulase having an enzymatic activity of at least 0.001 unit/mg solid [1 unit/mg solid forms 10μ mol of glucose from cellulose in one hour at 37°C at pH 8.3]. Also the amount of the alkali cellulase is such that enzymatic activity of alkali cellulase in the bath is preferably 0.1—1000 units/l, more particularly, 1—100 units/l.

Though the detergent composition of the present invention can be used in an unlimited pH range of from acidic to alkaline pH, it is preferred in order to sufficiently exhibit the deterging ffect of th alkali cellulase that the deterging bath is alkaline (more particularly, it has pH 7-11).

In the detergent composition of the present invention, components other than cellulase are not particularly limited. For example, the following components may be incorporated in the composition acc rding t their essential properties:

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Surfactants:

(1) Straight-chain or branched alkylbenzensulfonate salts having an alkyl group of 10—16 carbon atoms in average.

(2) Alkyl or alkenyl ether sulfate salts having a straight-chain or branched alkyl or alkenyl group of -20 carbon atoms in average, 0.5—8 mol in average of ethylene oxide, propylene oxide or butylene oxide in the molecule and an addition ratio of ethylene oxide/propylene oxide of 0.1/9.9-

9.9/0.1 or ethylene oxide/butylene oxide of 0.1/9.9—9.9/0.1. (3) Alkyl or alkenyl sulfate salts having an alkyl or alkenyl group of 10-20 carbon atoms in

everage. (4) Olefinsulfonate salts having 10—20 carbon atoms in average in the molecule.(5) Alkanesulfonate salts having 10—20 carbon atoms in average in the molecule.

(6) Saturated or unsaturated fatty acid salts having 10-24 carbon atoms in average in the

(7) Alkyl or alkenyl ether carboxylate salts having an alkyl or alkenyl group of 10-20 carbon

15 atoms in average, 0.5—8 mol in average of ethylene oxide, propylene oxide or butylene oxide in the molecule and an addition ratio of ethylene oxide/propylene oxide of 0.1/9.9-9.9/0.1 or ethylene oxide/butylene oxide of 0.1/9.9 to 9.9/0.1.

(8) α -Sulfo fatty acid salts or esters of the general formula:

20 wherein Y represents an alkyl group having 1—3 carbon atoms or a counter-ion, Z represents a 20 counter-ion and R represents an alkyl or alkenyl group having 10-20 carbon atoms.

As the counter-lons of anionic surfactants, there may be mentioned, for example, lons of alkali metals such as sodium and potassium, alkaline earth metals such as calcium and magnesium, ammonium, alkanolamines containing 1—3 alkanol groups having 2 or 3 carbon atoms such as monoethanolamine, diethanolamine, triethanolamine and trilsopropanolamine.

(9) Amino acid-type surfactants of the general formulae:

wherein R₁ represents an alkyl or alkenyl group having 8—24 carbon atoms, R₂ represents hydrogen or an alkyl group having 1-2 carbon atoms, R3 represents an amino acid residue and X represents an 30 alkali metal or alkaline earth metal ion.

No. 2
$$R_1$$
— CO — N — $(CH_2)_n$ — $COOX$ I R_2

wherein R_1 , R_2 and X have the same meanings as above and n represents an integer of 1—5.

wherein R_1 has the same meaning as above and m represents an integer of 1—8.

wherein R_1 , R_3 and X have the same meaning as above and R_4 r presents hydrogen, or an alkyl or hydroxyalkyl group having 1-2 carb n atoms.

wherein R_2 , R_3 and X have the same meaning as above and R_5 represents a β -hydroxyalkyl or β -40 hydroxyalkenyl group having 6-28 carbon atoms.

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wherein $R_{\mbox{\scriptsize 3}},\,R_{\mbox{\scriptsize 5}}$ and X have the same meaning as above.

(10) Phosphate ester surfactants:

No. 1 Acid alkyl (or alkenyl) phosphates:

wherein R' represents an alkyl or alkenyl group having 8-24 carbon atoms, n'+m' represents 3 and n' represents a number of 1-2.

No. 2 Alkyl (or alkenyl) phosphates:

10 wherein R' has the same meaning as above, n"+m" represents a number of 3 and n" represents a number of 1-3.

No. 3 Alkyl (or alkenyl) phosphate salts:

wherein R', n" and m" have the same meaning as above and M represents Na, K or Ca.

(11) Sulfonic acid-type amphoteric surfactants of the general formulae:

No. 1

wherein R₁₁ represents an alkyl or alkenyl group having 8-24 carbon atoms, R₁₂ represents an alkylene group having 1-4 carbon atoms, R₁₃ represents an alkyl group having 1-5 carbon atoms, R₁₄ represents an alkylene or hydroxyalkylene group having 1—4 carbon atoms.

20 No. 2

wherein R_{11} and R_{14} have the same meaning as above and R_{15} and R_{16} each represent an alkyl or alkenyl group having 8-24 or 1-5 carbon atoms.

No. 3

wherein R_{11} and R_{14} have the same meaning as above and n1 represents an integer of 1—20. 25

(12) Betaine-type, amphoteric surfactants of the general formulae:

No. 1

$$\begin{array}{c} R_{22} \\ | \\ R_{21} - N^{\odot} - R_{23} - COO^{\odot} \\ | \\ R_{22} \end{array}$$

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wherein R_{21} repr sents an alkyl, alkenyl, β -hydroxyalkyl or β -hydr xyalkenyl group having 8—24 carbon atoms, R₂₂ represents an alkyl group having 1—4 carbon atoms and R₂₃ represents an alkylene or hydroxyalkylene group having 1—6 carbon atoms.

$$(C_2H_4O)_{n2}H$$

 $|$
 R_{21} — N^{\oplus} — R_{23} — COO^{\oplus}
 $|$
 $(C_2H_4O)_{n2}H$

5 wherein R₂₁ and R₂₃ have the same meaning as above and n2 represents an integer of 1—20.

No. 3

wherein R_{21} and R_{23} have the same meaning as above and R_{24} represents a carboxyalkyl or hydroxyalkyl group having 2-5 carbon atoms.

(13) Polyoxyethylene alkyl or alkenyl ethers having an alkyl or alkenyl group of 10-20 carbon

10 atoms in average and 1—20 mol of ethylene oxide added.

(14) Polyoxyethylene alkylphenyl ethers having an alkyl group of 6—12 carbon atoms in average

and 1-20 mol of ethylene oxide added. (15) Polyoxypropylene alkyl or alkenyl ethers having an alkyl or alkenyl group of 10-20 carbon

atoms in average and 1-20 mol of propylene oxide added.

(16) Polyoxybutylene alkyl or alkenyl ethers having an alkyl or alkenyl group of 10-20 carbon atoms in average and 1-20 mol of butylene oxide added.

(17) Nonionic surfactants having an alkyl or alkenyl group of 10—20 carbon atoms in average and 1-30 mol in total of ethylene oxide and propylene oxide added or ethylene oxide and butylene oxide added (ratio of ethylene oxide to propylene oxide or butylene oxide being 0.1/9.9 to 9.9/0.1).

(18) Higher fatty acid alkanolamides or alkylene oxide adducts thereof of the general formula: 20

wherein R'11 represents an alkyl or alkenyl group having 10-20 carbon atoms, R'12 represents H or CH₃, n3 represents an integer of 1—3 and m3 represents an integer of 0—3.

(19) Sucrose/fatty acid esters comprising fatty acids having 10-20 carbon atoms in average 25

and sucrose. (20) Fatty acid/glycerol monoesters comprising fatty acids having 10-20 carbon atoms in average and glycerol.

(21) Alkylamine oxides of the general formula:

wherein R'₁₃ represents an alkyl r alkenyl group having 10—20 carbon atoms and R'₁₄ and R'₁₅ each 30 represent an alkyl group having 1-3 carbon atoms.

(22) Cationic surfactants f the gen ral formulae:

No. 1

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wherein at least one of R'_1 , R'_2 , R'_3 and R'_4 represents an alkyler alkenyl group having 8—24 carbon atoms and the remainder represents an alkyl group having 1—5 carbon atoms and X' represents a halogen.

wherein R'_1 , R'_2 , R'_3 and X' have the same meaning as above.

wherein R'₁, R'₂ and X' have the same meaning as above, R'₅ represents an alkylene group having 2—3 carbon atoms and n4 represents an integer of 1—20.

The composition preferably contains at least one of the above surfactants in an amount of at least 10 10 wt.%.

As preferred surfactants, there may be mentioned above surfactants 1), 2), 3), 4), 5), 6), 11)-No. 2, 12)-No. 1, 13), 14), 15), 17) and 18).

[2] Divalent metal ion sequestering agents:

The composition may contain 0—50 wt.% of one or more builder components selected from the group consisting of alkali metal salts or alkanolamine salts of the following compounds:

Salts of phosphoric acids such as orthophosphoric acid, pyrophosphoric acid, tripolyphosphoric acid, metaphosphoric acid, hexametaphosphoric acid and phytic acid.

2) Salts of phosphonic acids such as ethane-1,1-disphosphonic acid, ethane,1,1,2-triphosphonic acid, ethane-1-hydroxy-1,1-diphosphonic acid and its derivatives, ethane-hydroxy-1,1,2-triphosphonic acid, ethane-1-hydroxy-1,1,2-triphosphonic acid, ethane-1-hydroxy-1,1,2-triphosphonic acid, ethane-1-hydroxy-1,1,2-triphosphonic acid, ethane-1-hydroxy-1,1,2-triphosphonic acid, ethane-1-hydroxy-1,1,2-triphosphonic acid, ethane-1-hydroxy-1,1,2-triphosphonic acid, ethane-1-hydroxy-1,1,2-triphosphonic

acid, ethane-1,2-dicarboxy-1,2-diphosphonic acid and methanehydroxyphosphonic acid.
 3) Salts of phosphono carboxylic acids such as 2-phosphonobutane-1,2-dicarboxylic acids, 1-phosphonobutane-2,3,4-tricarboxylic acids and α-methylphosphonosuccinic acid.

4) Salts of amino acids such as aspartic acid, glutamic acid and glycine.

5) Salts of aminopolyacetic acids such as nitrilotriacetic acid, iminodiacetic acid, ethylenediaminetetracetic acid, diethylenetriaminepentaacetic acid, glycol ether diaminetetraacetic acid, hydroxyethyliminodiacetic acid, triethylenetetraminehexaacetic acid and dienkolic acid.

6) High-molecular electrolytes such as polyacrylic acid, polyaconitic acid, polyitaconic acid, polycitractonic acid, polyfumaric acid, polymaleic acid, polymesaconic acid, poly-α-hydroxyacrylic acid, polyvinylphosphonic acid sulphonated polymaleic acid, maleic anhydride/diisobutylene copolymer, maleic anhydride/styrene copolymer, maleic anhydride/methyl vinyl ether copolymer, maleic anhydride/ethylene copolymer, maleic anhydride/vinyl acetate copolymer, maleic anhydride/acrylonitrile copolymer, maleic anhydride/acrylate ester copolymer, maleic anhydride/butadiene copolymer, maleic anhydride/isoprene copolymer, poly-β-keto carboxylic acid derived from maleic anhydride and carbon monoxide, itaconic acid/ethylene copolymer, itaconic acid/aconitic acid copolymer, itaconic acid/maleic acid copolymer, itaconic acid/acrylic acid copolymer, maleic acid/maric acid copolymer, ethylene glycol/ethylene terephthalate copolymer, vinylpyrrolidone/vinyl acetate copolymer, 1-butene-2,3,4-tricarboxylic acid/itaconic acid/acrylic acid copolymer, quaternary ammonium group-containing polyester

polyaldehyde carboxylic acids, cis-isomer of epoxysuccinic acid, poly[N,N) bis(carboxymethyl)acrylamide], poly(hydroxy carboxylic acid), starch succinate, starch maleate, starch terephthalate, starch phosphate ester, dicarboxystarch, dicarboxymethylstarch and cellulose succinate esters.

7) Non-dissociating high-molecular compounds such as polyethylene glycol, polyvinyl alcohol, polyvinylpyrrolidone and cold water-soluble, urethanated polyvinyl alcohol.

8) Salts of dicarb xylic acids such as oxalic acid, malonic acid, succinic acid, glutaric acid, adipic acid, pimelic acid, suberic acid, azelalc acid and decane-1,10-dicarboxylic acid; salts of diglycolic acid, thiodiglycolic acid, xalacetic acid, hydroxydisuccinic acid, carboxymethylhydroxysuccinic acid and carboxymethyltartronic acid; salts of hydroxy carboxylic acids such as glycolic acid, malic acid, hydroxpivalic acid, tartaric acid, citric acid, lactic acid, gluconic acid, mucic acid, glucuronic acid and dialdehyde starch oxide; salts of itaconic acid, methylsuccinic acid 3-methylgiutaric acid, 2,2-dimethylmalonic acid, maleic acid, fumaric acid, glutamic acid, 1,2,3-propanetricarboxylic acid, aconitic acid, 3-butene-1,2,3-tricarboxylic acid, butane,1,2,3,4-tetracarboxylic acid,

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thanetetracarboxylic acid, ethenetetracarboxylic acid, n-alkenylaconitic acid, 1,2,3,4cyclopentanetetracarboxylic acid, phthalic acid, trimesic acid, hemimellitic acid, pyromellitic acid, benzenehexacarboxylic acid, tetrahydrofuran-1,2,3,4-tetracarboxylic acid and tetrahydrofuran-2,2,5,5tetracarboxylic acid; salts of sulfonated carboxylic acids such as sulfoitaconic acid, sulfotricarballylic acid, cysteic acid, sulfoacetic acid and sulfosuccinic acid; carboxymethylated sucrose, lactose and raffinose, carboxymethylated pentaerythritol, carboxymethylated gluconic acid, condensates of polyhydric alcohols or saccharides with maleic anhydride or succinic anhydride, condensates of hydroxy carboxylic acids with maleic anhydride or succinic anhydride, and organic acid salts such as CMOS and Builder M.

9) Aluminosilicates:

No. 1 Crystalline aluminosilicates of the formula:

 $\chi'(m'_2O \text{ or } M''O) \cdot Al_2O_3 \cdot \gamma'(SiO_2) \cdot w'(H_2O)$

wherein M' represents an alkali metal atom, M" represents an alkaline earth metal atom exchangeable with calcium, and x', y' and w' represent mole numbers of the respective components and generally, 15 they are as follows: 0.7≦x'≦1.5, 0.8≦y'≦6 and w' being a positive number.

No. 2 Detergent builders having the following general formula are particularly preferred:

Na₂O · Al₂O₃ · nSiO₂ · wH₂O

wherein n represents a number of 1.8—3.0 and w represents a number of 1—6. No. 3 Amorphous aluminosilicates of the formula:

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 $x(M_2O) \cdot Al_2O_3 \cdot y(SiO_2) \cdot w(H_2O)$

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wherein M represents sodium and/or potassium atom, and x, y and w represent mole numbers of the respective components within the following ranges:

0.7≦x≦1.2

1.6≦y≦2.8

w: any positive number including 0.

No. 4 Amorphous aluminosilicates of the formula:

$$X(M_2O) \cdot Ai_2O_3 \cdot Y(SiO_2) \cdot Z(P_2O_5) \cdot \omega(H_2O)$$

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wherein M represents Na or K and X, Y, Z and ω represent mole numbers of the respective components within the following ranges:

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0.20≤X≤1.10

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0.20≦Y≦4.00

0.001≦Z≦0.80

 ω : any positive number including 0.

[3] Alkalis or inorganic electrolytes:

The composition may contain also 1—50 wt.%, preferably 5—30 wt.%, of one or more alkali metal salts selected from the following compounds as the alkali or inorganic electrolyte: silicates, carbonates and sulfates. Further, the composition may contain organic alkalis such as triethanolamine, diethanolamine, monoethanolamine and triisopropanolamine.

[4] Antiredeposition agents:

The composition may contain 0.1—5% of one or more of the following c mpounds as antired position agent(s): polyethylene glycol, polyvinyl alcohol, polyvinylpyrr lidon and

carboxymethylcellulose. Particularly, a combination of carboxymethylcellulos or polyethylone glycol with the alkali cellulase of the present invention exhibits a synergism in the removal of muddy dirts.

In order to avoid the decomposition of carboxymethylcellulose by the alkali cellulase in the detergent composition, it is preferred that the carboxymethylcellulose is used in a granular or coated form.

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[5] BI aching agents:

A combination of the alkali cellulase of the present invention with a bleaching agent such as sodium percarb nate, s dium perb rate, sodium sulfate/hydrogen peroxide adduct or sodium chloride/hydrogen peroxide adduct or/and a photosensitive bleaching dye such as zinc or aluminum salt of sulfonated phthalocyanine further improves the deterging effects.

[6] Enzymes (enzymes which exhibits the essential enzymatic effects thereof in the deterging step):

As the enzymes, the following enzymes may be mentioned (classified with respect to their

enzymatic reactivities): Hydrolases, hydrases, oxido-reductases, desmolases, transferases and isomerases. All of these 10 enzymes may be used in the present invention. Particularly preferred enzymes are hydrolases such as proteases, esterases, carbohydrolases and nucleases.

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Examples of proteases are pepsin, trypsin, chymotrypsin, collagenase, keratinase, elastase, subtilisin, BPN, papain, bromelin, carboxypeptidases A and B, aminopeptidase and aspergillopeptidases

A and B. Examples of esterases are gastric lipase, pancreatic lipase, vegetable lipases, phospholipases,

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cholinesterases and phosphotases. Carbohydrolases other than alkali cellulases include maltase, saccharase, amylase, pectinase, lysozyme, lpha-glucosidase and eta-glucosidase.

[7] Bluing agents and fluoroescent dyes:

Various bluing agents and fluroescent dyes may be incorporated in the composition, if necessary. 20 For example, compounds of the following structures are recommended:

$$CH = SO_3N\alpha$$

$$CH = NH = NH$$

$$SO_3N\alpha$$

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and bluing agents f the general formulae:

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$$\begin{bmatrix}
D - NR - C & C & -Y \\
N & C & N
\end{bmatrix}$$
(S0₃H)₁

wherein D represents a residue of blue or purple, monoazo, disazo or anthraquinone dye, X and Y each represent hydroxyl group, amino group, an aliphatic amino group which may be substituted with hydroxyl, sulfonic acid, carboxylic acid or alkoxyl group, or an aromatic or alicyclic amino group which 5 may be substituted with a halogen atom or hydroxyl, sulfonic acid, carboxylic acid, lower alkyl or lower alkoxyl group, R represents a hydrogen atom or a lower alkyl group but excluding cases wherein (1) R represents a hydrogen atom and both X and Y represent a hydroxyl group or an alkanolamine at the same time and (2) R represents a hydrogen atom, one of X and Y represents a hydoxyl group and the other represents an alkanolamine group, and n represents an integer of at least 2, and

$$D-NH-C N C-X$$

$$N C N$$
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wherein D represents a residue of a blue or purple, azo or anthraquinone dye, and X and Y may be the same or different and represent an alkanolamine residue or a hydroxyl group.

[8] Caking-preventing agents:

The following caking-preventing agents may be incorporated in powdery detergent composition: 15 p-toluenesulfonate salts, xylenesulfonate salts, acetate salts, sulfosuccinate salts, talc, finly pulverized 15 silica, clay, calcium silicates (such as Micro-cell of Johns-Manvill Co.), calcium carbonate and magnesium oxide.

[9] Masking agents for factors inhibiting the alkali cellulase activity:

The alkali cellulases are deactivated in some cases in the presence of copper, zinc, chromium, 20 mercury, lead, manganese or silver ions or their compounds. Various metal chelating agents and metalprecipitating agents are effective on these inhibitors. They include, for example, divalent metal ion sequestering agents as listed in the above item [2] with reference to optional additives as well as magnesium silicate and magnesium sulfate.

Cellobiose, glucose and gluconolactone act sometimes as the inhibitors. It is preferred to avoid the 25 co-presence of those saccharides with the alkali cellulase as far as possible. In case the co-presence is unavoidable, it is necessary to avoid the direct contact of the saccharides with the alkali cellulase by, for example, coating them.

Strong chelating agents such as ethylenediaminetetraacetate salts, anionic surfactants and cationic surfactants act as the inhibitors in some cases. However, the co-presence of those substances with the alkali cellulase is allowable if the direct contact of them is prevented by some ideas such as 30 tableting and coating methods.

The above-mentioned masking agents and methods may be employed, if necessary, in the present invention.

[10] Alkali cellulase-activators:

The activators vary depending on variety of the alkali cellulases. In the presence of proteins, cobalt and its salts, calcium and its salts, potassium and its salts, sodium and its salts or monosaccharides such as mann se and xyl se, the alkali cellulases ar activated and their deterging powers ar impr ved remarkably.

[11] Antioxidants:

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The antioxidants includ , for example, tert-butylhydroxytoluene, 4,4'-butylidenebis(6-tert-butyl-3-methylphenol), 2,2'-butylidenebis(6-tert-butyl-4-methylphenol), monostyrenated cresol, distyrenated cresol, monostyrenated phenol, distyr nated phenol and 1,1'-bis(4hydroxyphenyl)cyclohexane.

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[12] Solubilizers:

The solubilizers include, f r example, lower alcohols such as ethanol, benz nesulfonate salts, lower alkylbenzenesulfonate salts such as p-toluenesulfonate salts, glycols such as pr pylen glycol, acetylbenzenesulf nate salts, acetamides, pyridinedicarboxylic acid amides, benzoate salts and urea.

The following examples will further illustrate the present invention. In the following referential example, the preparation of an alkali cellulase is explained. Unless otherwise state, percentages in the following examples are given by weight.

Referential Example 1

Preparation of alkali cellulase:

Alkali-resistant cellulases according to the present invention are obtained by, for example, a technique disclosed in G. Okada, T. Nishizawa and K. Nishizawa "Biochem. J., 99, 214 (1966)". More particularly, a crude enzyme solution was extracted from the hepatopancreas of a marine mollusc (Dolabella sp.). The crude enzyme solution was subjected to the starch zone-electrophoresis and carboxymethylcellulose-saccharifying activity of the resulting fraction was measured. The 15 carboxymethylcellulose-saccharifying activity of pH 8.3 was determined from an absorbance (ΔΟD) at 660 m μ using an alkaline copper reagent and arsenomolybdate after reacting the fraction with carboxymethylcellulose.

Cellulase Activity at pH 8.3 (AOD) Fraction No. 20 0.05 10 0.55 15 0.47 20 0.40 30 0.12 35

It is understood that fractions Nos. 15, 20 and 30 contain cellulases having a high activity under 25 25 a weak alkaline condition.

Example 1

Effects of alkali cellulase superior to those of other enzymes on cotton cloths artificially stained with muddy dirts will be shown:

	41	The second secon	
วก	7)	Detergent compositions:	

, ,	Detergent compositions.			•
		A (%)	B (%)	
	Sodium straight-chain dodecylben- zenesulfonate	10	.—	
35	Sodium α -olefinsulfonates (C_{16-18})	5 2		35
	Sodium alkylethoxysulfates (C ₁₄₋₁₅ , EO=1.5 mols)	2	25	
	Sodium alkyl sulfates (C ₁₄₋₁₅)	3	_	
	Soap (beef fatty acid sodium salts)	2	_	
40	Secondary alcohol (C=13.5) ethoxylate (EO=7)		25	40
	Sodium tripolyphosphate	10		
	Crystalline sodium aluminosilicate (type 4A)	10	_	
45	Sodium silicate	10		45
	Triethanolamine		5	
	Sodium carbonate	10	-	
	Potassium carbonate	_	5	
	Carboxymethylcellulose	1	1	
50	Polyethylene glycol (MW) 6000)	1	1	50
00	Fluorescent dye	0.4	0.3	
	Bluing agent	<u>-</u>	0.05	
	Sodium p-toluenesulfonate	2		
	Ethanol		8	
55	Water	. 10	balance	55
55	Enzyme	0 r3	0 or 2	
	Perfume	0.2	0.1	
	Glauber's salt	balance		

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2) Mud-stained loths (artificially stained cloths):

Kanuma s kigyoku soil for horicultural use was dried at 120°C±5°C for 4 h and then pulverized. 150 mesh (100 μ m)-passed soil particles were dried at 120°C \pm 5°C for 2 h. 150 g of the soil particles was dispersed in 1000 I of Perclene. A calico #2023 cloth was contacted with the dispersion and 5 brushed. After removal of the dispersion, excessive mud remaining on the cloth was removed (Japanese Patent Laid-Open No. 26473/1980).

Test pieces having a size of 10 cm x 10 cm were prepared and subjected to the tests.

3) Deterging conditions and method and appraisal:

A detergent was dissolved in 4°DH hard water to obtain 1 l of 0.133% aqueous detergent 10 solution. Five pieces of the cloth artificially stained with the muddy dirt were immersed in the aqueous detergent solution. After leaving them to stand at 40°C for 2 hours, the detergent solution and the pieces of artificially stained cloth were transferred in a stainless steel beaker for Turgotometer and stirred at 100 rpm at 20°C for 10 minutes in the Turgotometer. After washing with running water, they were pressed with an iron and their reflectivities were measured. The deterging rate was calculated 15 according to the following formula:

Reflectivities of the original cloth before the washing and those of the stained cloth before and after the washing were measured by means of an automatic recording colorimeter (a product of Shimadzu Seisaku-sho) and the deterging rate (%) was calculated according to the following formula:

	(Reflectivity after washing)		(Reflectivity before washing)	×100
Deterging rate (%)=-	(Reflectivity of original cloth)	_	(Reflectivity before washing)	

An average of fives samples was shown in Table 1. 20 The aqueous detergent solution before the washing had pH 10.6. 20

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4) Enzymes used:

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40 5) Results:

(1) Not used (balanced with Glauber's salt) (comparative)

(2) Alkali cellulase (crude enzyme solution of fraction No. 15 in the above 25 referential example) (3) Alkali cellulase (crude enzyme solution Present of fraction No. 20 in the above invention

referential example) (4) Alkali cellulase (crude enzyme solution of fraction No. 30 in the above

referential example) (5) Cellulase (crude enzyme solution of fraction No. 35 in the above referential example)

(6) Amylase (Termamyl 60 G; a product of Novo Co.)

(7) Protease (Gist Brocades, maxatase P 330,000) (8) Lipase (Olipase; a product of Nagase Sangyo Co.)

Table 1

		Deterging rate (%)		
	Enzyme in the Detergent	Composition A	Composition B	
45	① ②	65 81	60 78	45
	3 •	80 79	77 76	
50	6 6	74 66	72 —	50
50	0 ®	65 65	_	•

		·	
	5	A culture medium (having a pH value f 10) c ntaining 1.0% f peptone, 1.0% of meat extract, 1.0% of carb xymethyl cellulose (CMC), 0.5% of sodium chloride, 0.1% of potassium dihydrogen phosphate and 1.0% of anhydrous sodium carbonate was inoculated with <i>Bacillus</i> N4, a novel species belonging to the genus <i>Bacillus</i> (deposited with the FRI deposition number of 1141), separated from the soil collected at Hirosawa, Wako city, Saitama prefecture, and shaking culturing was carried out 37°C for 72 hours. Cells were removed by centrifugal separation to obtain a crude enzyme. The crude enzyme was dried with ethanol according to the customary method to obtain a cellulase powder. Thus, 10 g/l of a cellulase enzyme (having an enzymatic activity of 0.6 unit/mg of the soild at a pH value of 6) (hereinafter referred to as "cellulase N-4") was obtained. At a pH value of 9, the so-obtained enzyme retained 85% of the activity at a pH value of 6. Incidentally, a commercially available cellulase originating from <i>Aspergillus niger</i> had an activity of 0% at a pH value of 9. That is, the cellulase had no activity at a pH value of 9.	5
1	15	Referential Example 3 A flask was charged with 9 ml of a culture medium containing 0.5% of ammonium sulfate, 1.5% of pulp block, 0.02% of glucose, 0.1% of yeast extract, 0.02% of MgSO₄ · 7H₂O and 0.2% of K₂HPO, and the culture medium was sterilized at 120°C for 20 minutes. The sterilized culture medium was cooled	15
:	20	and mixed with 10 ml of 0.7% aqueous solution of NaHCO ₃ separately sterilized. The cuttle medium was then inoculated with a cellulase 212-producing species belonging to the genus Aeromonas (deposited with the FRI deposition number of 2306), and shaking culturing was carried out at 37°C for 72 hours. Cells were removed by centrifugal separation to obtain a crude enzyme liquid of cellulase 212. The crude enzyme liquid was dried with ethanol according to the customary procedure to obtain a 212-medium procedure to obtain a crude enzyme liquid was dried with ethanol according to the customary procedure to obtain a 212-medium procedure to obtain a 212-medi	20
;	25	(hereinafter referred to as "cellulase 212"). At a pH value of 9, the so-obtained enzyme retained 70% of the enzymatic activity at a pH value of 6. Enzymes used in Examples 2 to 7 are listed as follows. (1) Cellulase N4	25
	30	 (2) Cellulase 212 (3) Cellulase (supplied by Sigma Co., originating from Aspergillus niger, 1.35 units/mg) (4) Lipase (supplied by Gist Brochades NV, originating from R. oryzae) (5) Amylase (Termamil 60G supplied by Novo Industries Co.) (6) Protease (Alkalase 2.0M supplied by Novo Industries Co.) 	30
	35	Example 2 A highly alkaline powdery detergent for clothing was prepared according to the following recipe. The pH value of a 0.133% aqueous solution of the detergent was 11.2.	35
		Sodium linear-dodecylbenzene-sulfonate Soap (sodium salt of beef tallow fatty acid) Solium satton bosphate 20% by weight 20% by weight	
	40	Sodium orthophosphate Sodium metaphosphate Sodium carbonate Carboxymethyl cellulose Polyothylogo glycol 1% by weight 1% by weight 1% by weight	40
	45	Polyethylene glycol Fluorescent dye Glauber salt Enzyme O.4% by weight balance O or 2% by weight 5% by weight	45
	50	The results of the washing test made on so-prepared detergents are shown in Table 2. Incidentally, in Table 2 and Tables given hereinafter, each detergent is identified by example number—(0)).	50
		Table 2	
		Washing Detergent Power Index	
	55	1—(2) (present inv_ntion) 104.0 1—(3) 101 1—(4) 100	55
	_	1—(5)	60
	60) 1—(6)	

.3				
	Exampl 3	detergent for clothing Wi	is prepared accirding to the fillowing recipe.	
	A weakly alkali	ne powdery detergent for clothing we	ent was 10.3.	
	The pH value of a 0.1	33% aqueous solution of the deterge	ME 1743 1010.	
	•			
		Sodium $lpha$ -olefin-sulfonate	20% by weight	5
5		Soap	1% by weight	•
•		Sodium tripolyphosphate	20% by weight	
		Sodium silicate (JIS No. 2)	10% by weight	
		Sodium carbonate	5% by weight	
		Sodium carbonate	1% by weight	
		Carboxymethyl cellulose	1% by weight	10
10		Polyethylene glycol	0.4% by weight	
		Fluorescent dye		
		Glauber salt	balance	
		Enzyme	0 or 2% by weight	
		Water	10% by weight	
		7.77		
		described in Evennia 1, the	washing test was carried out. The obtained	15
15	In the same ma	inner as described in Example 1, the	washing toot was assistance	
	results are shown in	Table 3.		
	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,			
		Table 3		
			Washing Power	
		Detergent	Index	
		Detergent		
			100	20
20		2—(0) (reference detergent)	104	
20		2-(1) (present invention)		
	•	2—(2) (present invention)	104.5	
			100	
		2—(4)	100	
		2(5)	100.5	25
25		2—(6)	100.0	
20				
	Example 4			
	A soutral now	tery detergent for clothing was prepa	red according to the following recipe. The pH	
	A figural power	queous solution of the detergent was	6.8	•
	value of a U. 13376 at			
		Sodium linear-alcohol (C=14) sulfa	ate 30% by weight	-00
		Sodium linear-alcohol (C-147 out)	1% by weight	30
30		Polyethylene glycol		
•-		Sodium phosphate	1% by weight	
		Fluorescent dye	0.2% by weight	
		Glauber salt	balance	
			0 or 2% by weight	0.5
		Enzyme	5% by weight	35
35		Water	determents are shown in Table 4.	
	The results of	Water the washing test made on so-prepare	ad deferdents are shown in the	
	1110 1000111		•	
	•	Table 4		
			Washing	
		Detergent	Power Index	
		Decetyett		
		- 101 / C	100 ·	40
40	•	3—(0) (reference detergent)	103	
40		3(1) (present invention)		
		3—(2) (present invention	103.5	
		3—(4)	· 100	
			100	
	•	3—(5)	100	45
45		3—(6)	•	
70				
	Example 5		ding to the following recipe.	
	A shoenhorous	s-free, weakly alkaline detergent was	prepared according to the following recipe.	
	W hinshinger			
		Sodium linear-dodecylbenzene-	15% by weight	
		Social inigat-doggoding in		
		sulfonate	5% by weight	50
50		S dium alkylethoxy-sulfate	Over Holding	
		(C ₁₄ —C ₁₅ , EO—3 m les)	a and become be	
		Builder and enzyme	20% by weight	•
		(see Tabl 5)		
		Sodium silicate	15% by weight	
				==
		Sodium silicate		55
55		Sodium carbonate	15% by weight	99

•					
4				GB 2 095 275 A	14
	Carboxymethyl cellulos Polyethylene glycol Fluorescent dye Glauber salt	se	1.5% by weight 1.5% by weight 0.5% by weight balance		r
5	Water The results of the washing test are show	wn in Tablo Table !		,	5
	Builder Enz	ryme	Washing Power Index		
10	sodium tripolyphosphate, 20% — sodium citrate, 20% — zeolite type 4A, 20%	98 98	.5	t)	10
15	zeolite type 4A, 15% (6), sodium citrate, 15% (1), sodium citrate, 15% (2), zeolite type 4A, 15% (1),	, 5% 10 , 5% 10			15
20	Example 6 Detergents were prepared according to enzymes. The results of the washing test man	the recipe de on thes	adopted in Example 3 e detergents are show	by using combinations of n in Table 6.	20
		Table (S nbination of Enzymes		
	Detergent		ht Number Indicates % of Enzyme)	Washing Power Index	
25	2—(2) (reference detergent) 2—(2)/(4) (present invention 2—(2)/(5) (present invention	(2)/(l)=1/1 5)=1/1	100 100.5 100.5 101	25
30	2—(2)/(6) (present invention) 2—(2)/(4)/(6) (present invention) 2—(2)/(5)/(6) (present invention 2—(4)/(5)/(6)	(2)/(4 (2)/(5	5)=1/1 1}/(6)=2/1/1 5]/(6)=2/1/1 5]/(6)=2/1/1	101.5 101.5 98	30
	Example 7 A weakly alkaline powdery detergent for	or clothing	was prepared according	ng to the following recipe.	
35	Sodium alkyl-sulfate (C= Sodium alkylethoxy-sulfa (C=14.5, EO=3)	14.5)	15% by weigh 5% by weight	nt	35
40	Soap (beef tallow type) Sodium pyrophosphate Sodium silicate Sodium carbonate		2% by weight 18% by weigh 13% by weigh 5% by weight 2% by weight	nt nt	40
45	Polyethylene glycol Fluorescent dye Glauber salt Magnesium silicate Water Enzyme		0.2% by weight balance 1% by weight 5% by weight 2% by weight	ht	45
	Sodium percarbonate		15% by weigh		
	The results of the washing test made o	n so-prepa Tabi)	
50	Detergent	Enzyme	, Washing Power Index		50
	6—(6) (reference detergent) 6—(1) (present invention) 6—(2) (present invention)	(6) (1) (2)	100 102.5 103	-	

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-		•	
4	9	12	าร
•	a		13

1. A deterg nt composition which includes an alkali cellulase.

2. A detergent composition as claimed in Claim 1, in which said alkali cellulase has an optimum effectiveness at a pH of 8.0 to 11.5.

3. A detergent composition as claimed in Claim 1, in which said alkali cellulase is produced by a Bacillus N fungus or a cellulase 212-producing fungus belonging to the genus Aeromonas.

4. A detergent composition as claimed in Claim 1, which contains 0.01 to 70 wt.% of said alkali cellulase having an enzymatic activity of at least 0.001 unit/mg solid.

5. A detergent composition as claimed in Claim 1, which contains said alkali cellulase in an

10 amount of 0.1 to 1000 units per liter of the composition. 6. A detergent composition as claimed in Claim 1, which further comprises one or more of the following surfactants, divalent metal ion sequestering agents, alkali agents, inorganic electrolytes, antiredeposition agents, bleaching agents, enzymes, bluing agents, fluorescent dyes, caking-preventing agents, masking agents for factors which inhibit the alkali cellulase activity, activators for the alkali cellulase, antioxidants, solubilizers and other conventional additives.

7. A detergent composition as claimed in Claim 1, in which said alkali cellulase is extracted from the hepatopancreas of a marine mollusc.

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